

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 808 860 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

26.11.1997 Bulletin 1997/48

(51) Int. Cl.⁶: C08G 61/12

(X)

(21) Application number: 97108237.5

(22) Date of filing: 21.05.1997

(84) Designated Contracting States:

DE FR GB

- Aoki, Kei
- Ikoma-shi, Nara-ken (JP)

(30) Priority: 22.05.1996 JP 151546/96

(71) Applicant:

NIPPON PAINT CO., LTD.
Kita-ku, Osaka (JP)

(74) Representative:

Kinzebach, Werner, Dr. et al
Patentanwälte
Reitstötter, Kinzebach und Partner
Postfach 86 06 49
81633 München (DE)

(72) Inventors:

- Irie, Takashi
Sulta-shi, Osaka-fu (JP)

(54) Curable resin composition for coating use

(57) A resin composition curable through a Michael reaction between (a) a component having a plurality of α, β -unsaturated carbonyl groups and (b) a component having a plurality of activated methylene group in the presence of a catalyst is disclosed. The activated methylene component is a polymer of an asymmetric malonate ester in which one of carboxyl groups is esterified with an alkanol while the other carboxyl group is esterified with hydroxylalkyl (meth) acrylate or polyoxyalkylene glycol mono(meth)acrylate.

EP 0 808 860 A2

Description**FIELD OF THE INVENTION**

5 This invention relates to a curable resin composition for coating use. More particularly, it relates to a resinous composition which cures through a Michael reaction.

BACKGROUND OF THE INVENTION

10 Michael reactions in which base-catalysed addition of activated methylene to polarized double bond are utilized in the crosslinking of resinous compositions for coating use. These compositions are advantageous in that the Michael reaction gives a chemically stable crosslinking bond without forming any reaction by-product.

Typical examples of prior art include the following patent literature:

15 U.S. Patent No. 4,408,018;
 U.S. Patent No. 4,602,061;
 U.S. Patent No. 5,017,649;
 U.S. Patent No. 5,084,536;
 EP-A-0448154;
 20 JP-A-01/121341 (EP-A-0310011); and
 JP-A-01/204919 (EP-A-0326723).

Typical examples of activated methylene compounds used in the Michael reaction are derivatives of acetoacetic acid, cyanoacetic acid and malonic acid. For use in the crosslinking reaction of resins or polymers, the Michael donor 25 must be a compound or polymer having a plurality of activated methylene groups in the molecule. Acrylic monomers having an acetoacetoxy group can be synthesized relatively easily by reacting diketene with a hydroxylalkyl acrylate or methacrylate such as 2-hydroxyethyl acrylate (HEA) or 2-hydroxyethyl methacrylate (HEMA). Activated methylene-containing acrylic polymers used in the prior art are either polymers of acetoacetylated hydroxylalkyl (meth)acrylate monomers or acetoacetylated acrylic polyols produced by reacting diketene with an acrylic polyol polymer.

30 Because malonic acid is a dibasic acid, malonate groups can be incorporated into a polyester chain to produce a polymeric Michael donor. U.S. Patent No. 4,602,061 discloses a liquid two-component coating composition comprising (a) an oligomeric or polymeric malonate compound such as polyurethanes, polyesters, polyacrylates, epoxy resins, polyamids or polyvinyl resins containing malonate groups, either in the main chain or the side chain or in both, (b) an α , β -ethylenically unsaturated carbonyl compound and (c) a basic catalyst.

35 The malonate group-containing polyacrylate resin disclosed therein is a reaction product of the above-mentioned malonate polyester resin first with a diisocyanate and then with HEA or HEMA.

JP-A-03206012 discloses a malonate group-containing acrylic monomer and polymers thereof for use as odorless, low irritating and low water absorbing dental materials. Corresponding acrylate monomers and polymers containing higher homologs such as succinate or adipate are also disclosed. The use of these monomers and polymers in combination with a Michael acceptor and a catalyst are not disclosed therein.

40 It has been discovered that resinous compositions containing a malonate group-containing acrylate polymer, a Michael acceptor and a catalyst are advantageous in terms of weatherability over corresponding compositions containing an acetoacetate group-containing acrylate polymer when used for coating purposes.

45 SUMMARY OF THE INVENTION

According to the present invention, a curable resin composition having enhanced weatherability is provided by using as an activated methylene component a malonate group-containing acrylate polymer in place of an acetoacetate group-containing acrylate polymer.

50 Accordingly, the present invention provides a curable resin composition comprising:

- (a) a component containing a plurality of α , β -ethylenically unsaturated carbonyl groups in the molecule;
- (b) an acrylate polymer containing a plurality of malonate-terminated pendant groups in the molecule; and
- (c) a catalyst capable of promoting the Michael reaction.

55 The present invention also provides a coating composition containing the above resin composition as a vehicle resin.

DESCRIPTION OF THE PREFERRED EMBODIMENTSComponent (a)

5 Component (a) is a compound or polymer having a plurality of ethylenic unsaturations between carbon atoms at the α and β positions relative to a carbonyl group. Typical examples of such compound are acrylic or methacrylic (hereinafter collectively referred to as "(meth) acrylic") esters of polyhydric alcohols such as ethylene glycol di(meth)acrylate, diethylene glycol di(meth)acrylate, propylene glycol di(meth)acrylate, neopentyl glycol di(meth)acrylate, trimethylolpropane tri(meth) acrylate, glycerol tri(meth)acrylate, pentaerythritol tetra(meth)acrylate and the like.

10 Unsaturated polyesters containing an unsaturated dicarboxylic acid such as maleic or fumaric acid may also be used as component (a). Unsaturated polyesters are used in large quantities for molding various FRP products and may be synthesized, as is well-known, by the polycondensation reaction between an α , β - unsaturated dicarboxylic acid component such as maleic anhydride or fumaric acid optionally containing other polycarboxylic acids such as phthalic anhydride, isophthalic, terephthalic, adipic, sebacic or trimellitic acid and a polyol component such as ethylene glycol, diethylene glycol, propylene glycol, tetramethylene glycol, 1,6-hexanediol, neopentyl glycol, trimethylolpropane, glycerine or pentaerythritol.

15 Other examples of component (a) include epoxy acrylate resins produced by reacting (meth)acrylic acid with an epoxy resin such as bisphenol or novolac type epoxy resins, and urethane acrylate resins produced by reacting a hydroxyl group-containing acrylic resin first with an organic polyisocyanate such as isophorone diisocyanate, tolylene diisocyanate, diphenylmethane diisocyanate, hexamethylene diisocyanate or a urethane prepolymer thereof and then with 2-hydroxyethyl (meth)acrylate.

20 Acrylic resins containing a plurality of α , β - unsaturated carbonyl functions may be produced via the ring-opening reaction of an epoxy ring with (meth)acrylic acid or the acylation of a hydroxyl group with (meth)acrylic acid. For example, epoxy group-containing acrylic resins may be produced by copolymerising an epoxy group-containing acrylic monomer such as glycidyl (meth)acrylate with an acrylic monomer free of the epoxy group and/or a nonacrylic ethylenically unsaturated monomer. Examples of copolymerizable acrylic monomers include alkyl (meth) acrylates such as methyl, ethyl, propyl, n-butyl, isobutyl, t-butyl, 2-ethylhexyl or lauryl (meth)acrylate; hydroxyalkyl (meth)acrylates such as 2-hydroxyethyl (meth) acrylate; aryl or aralkyl (meth)acrylates such as phenyl or benzyl (meth)acrylate; PLACCEL FA1 or FM1 (adduct of 2-hydroxyethyl (meth)acrylate with polycaprolactone sold by Daicel Chemical Industries, Ltd.); and 25 other acrylic monomers such as acrylamide, methylene-bis-acrylamide or acrylonitrile. Examples of copolymerizable non-acrylic monomers include styrene, α -methylstyrene, itaconic acid, maleic acid, vinyl acetate and the like. The epoxide function of the acrylic copolymer may then be opened with (meth)acrylic acid to introduce the α , β - unsaturated carbonyl function. Conversely, an epoxy group-containing acrylic monomer such as glycidyl (meth)acrylate may be reacted with an acrylic polymer containing a plurality of free carboxylic functions which may be produced by copolymerizing a carboxyl group-containing monomer such as acrylic, methacrylic or itaconic acid with a copolymerizable acrylic monomer and optionally with a non-acrylic monomer as discussed supra.

30 Acryl polyols containing a plurality of hydroxyl groups may be modified to have a plurality of α , β - unsaturated carbonyl functions by the acylation reaction with acrylic or methacrylic acid or a reactive derivative thereof such as chloride, or the transesterification reaction with a lower alkyl (meth)acrylate such as methyl (meth)acrylate. Acryl polyols may be produced by copolymerizing a hydroxyl group-containing acrylic monomer such as 2-hydroxyethyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, 4-hydroxybutyl (meth)acrylate or PLACCEL FM-1 with a copolymerizable acrylic and/or non-acrylic monomer as discussed supra.

35 Further examples of component (a) include polyether acrylate resins such as polyethylene glycol di(meth) acrylate, silicone oligomers such as 3-methacryloyloxypropyl-terminated polydiorganosiloxanes and fluororesins containing α , β - unsaturated carbonyl groups.

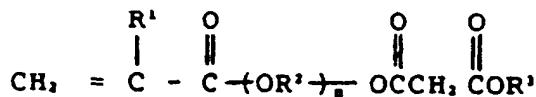
40 When component (a) is a resin, its molecular weight ranges between 400 and 100,000, preferably between 600 and 10,000, and the alkenyl equivalent weight ranges between 100 and 10,000, preferably between 100 and 1,000. The term "equivalent weight" as used herein refers to the molecular weight divided by the number of a function concerned.

Component (b)

45 Component (b) may be produced by copolymerizing a malonate-terminated acrylate monomer with a copolymerizable acrylic and/or non-acrylic monomer as exemplified in connection with Component (a).

50 Usable malonate-terminated acrylate monomers include but are not limited to the acrylate monomer disclosed in JP-A-03206012 cited supra. The malonate-terminated acrylate monomers have the formula:

5



wherein R¹ is hydrogen or methyl, R² is a C₂-C₆-alkylene, R³ is a C₁-C₆ alkyl and n is an integer from 1 to 10.

10 A first class of preferred monomers are monomers of the above formula wherein R² is -CH₂CH₂-, n is 1 and R³ is a C₁-C₆ alkyl. These monomers are produced by the transesterification reaction of dialkyl malonate with HEA or HEMA.

A further class of preferred monomers are produced by transesterifying dialkyl malonate with a polyoxyalkylene glycol monoacrylate or methacrylate. Monomers of this class are represented by the above formula wherein R² is -CH₂CH₂-, -CH(CH₃)CH₂- or -(CH₂)₄, n is an integer of greater than 1, and R³ is a C₁-C₆ alkyl.

15 Examples of copolymerizable acrylic monomers include alkyl (meth)acrylates such as methyl, ethyl, propyl, n-butyl, isobutyl, t-butyl, 2-ethylhexyl or lauryl (meth)acrylate; hydroxyalkyl (meth)acrylates such as 2-hydroxyethyl (meth) acrylate; aryl or aralkyl (meth)acrylates such as phenyl or benzyl (meth)acrylate; and other acrylic monomers such as acrylamide, methylene-bis-acrylamide or acrylonitrile. Examples of copolymerizable non-acrylic monomers include styrene, α-methylstyrene, itaconic acid, maleic acid, vinyl acetate and the like.

20 Component (b), namely acrylate polymers having a plurality of malonate-terminated pendant groups preferably have a number average molecular weight from 1,000 to 50,000, more preferably from 1,000 to 10,000, and an activated methylene equivalent weight from 100 to 10,000, more preferably from 150 to 5,000.

Component (c)

25 As is well known, the Michael reaction normally requires the presence of a strong base to strengthen the acidity of methylene proton by the conversion of adjacent carbonyl group to an enolate anion.

30 Component (c) of the resin composition of the present invention may be a strong base. Examples thereof include alkali metal hydroxide such as sodium hydroxide or potassium hydroxide; alkali metal alkoxide such as sodium methoxide or potassium ethoxide; quaternary ammonium hydroxides such as tetrabutylammonium hydroxide or benzyltrimethylammonium hydroxide; quaternary ammonium carbonates such as tetrabutylammonium carbonate or benzyltrimethylammonium carbonate; tertiary amines such as diazabicyclooctane otherwise named DABCO; quanidines and amidines such as tetramethylquanidine, 1,8-diazabicyclo[5.4.0]undecene-7 or diazabicyclo[4.3.0]nonene-5; tertiary phosphines such as triphenylphosphine; and a mixture thereof. Quaternary ammonium fluorides and tetrahydroborates such as tetrabutylammonium fluoride or benzyltrimethylammonium fluoride or corresponding tetrahydroborates exhibit a catalytic activity by themselves in the Michael reaction as same as the above-mentioned strong bases.

35 Be cause strong bases and quaternary ammonium fluorides or tetrahydroborates are capable of catalyzing the Michael reaction even at room temperature, they cannot be incorporated into a storage stable, one-component curable resin composition. When such a one-component composition is desired, the strong base may be blocked with a carboxylic acid which undergoes volatilization or decarboxylation under baking conditions. Storage stable, one-component compositions may also be formulated by the use of a combination of an onium salt having no or little catalytic activity and an epoxy compound. The epoxy compound may be added to the composition as a discrete component or it may be covalently attached to either Component (a) or Component (b). This catalyst system is disclosed in commonly assigned U.S. Patent No. 5,565,525 of which disclosure is incorporated herein by reference.

40 A variety of ammonium salts may be used. However, tetraalkyl and trialkylaralkyl type ammonium salts are preferred for economical reasons. Nitrogen-containing heterocyclic ammonium salts such as those derived from pyridine, piperidine, piperazine or morpholine may also be used. Specific examples of ammonium cations include tetrabutylammonium, tetramethylammonium, tetraethylammonium, trimethylbenzylammonium, tetrapropylammonium, tetrahexylammonium, tetraoctylammonium, tetradecylammonium, tetrahexadecylammonium, triethylhexylammonium, 2-hydroxyethyltrimethylammonium (choline), methyltriocetylammmonium, cetyltrimethylammonium, 2-chloroethyltrimethylammonium and methylpyridinium.

45 Counter anions are selected to form a non-basic stable salt with the ammonium cations and include halides other than fluoride, carboxylates, sulfonates, nitrate, sulfate, sulfite, phosphate and the like. Specific examples thereof include acetate, laurate, glycolate, benzoate, salicylate, maleate, phthalate, chloride, bromide, iodide, methanesulfonate, p-toluenesulfonate, dodecylbenzene sulfonate, triflate, nitrate, sulfate, methosulfate, phosphate and acid t-dibutylphosphate. Quaternary phosphonium salts such as tetrabutylphosphonium bromide and tertiary sulfonium salts such as trimethylsulfonium iodide may also be used.

50 Onium salts, among others, having counter anions derived from phenol carboxylic acids such as salicylic acid, polybasic carboxylic acid such as oxalic, maleic, fumaric or phthalic acid, nitric acid, phosphoric acid, sulfuric acid, phos-

phate ester or sulfonic acid exhibit a catalytic activity only in the presence of an epoxy component under the conventional baking condition. Accordingly, they are useful when a thermosetting, storage stable, one-component composition is desired.

In certain applications where the curable composition of the present invention is applied onto a pigmented base coat wet-on-wet and baked simultaneously with the base coat as in car body finishing, the catalyst can diffuse into the base coat layer so that the composition is not fully cured. Accordingly, it is advantageous to immobilize the catalyst in component (a) or component (b). This may be accomplished by the use of ammonium salts having a functional group capable of linking to component (a) or component (b). When component (a) or component (b) has a free carboxyl group, glycidyl group-containing ammonium salts such as glycidyltrimethylammonium chloride or bromide will react with the carboxyl group to form a covalent linkage. Such an ammonium salt may be added as a discrete component or as a conjugate with component (a) or component (b). Quaternary ammonium salts having acryloyl or methacryloyl group such as 2-(methacryloyloxy)-ethyltrimethylammonium chloride or bromide may also be used for this purpose. These ammonium monomers may be copolymerized with other monomers when producing component (a) or component (b). When added as a discrete component to the resin composition, they react with component (a) as a Michael donor.

When used as a discrete component, preferred examples of epoxy compounds include glycidyl ethers such as phenyl glycidyl ether, bisphenol epoxy resins, reaction products of epichlorohydrine with a polyhydric alcohol such as ethylene glycol or trimethylolpropane; glycidyl esters such as glycidyl benzoate, glycidyl (meth)acrylate or its polymers; alicyclic epoxides such as 4-(3,4-epoxycyclohexyl) methoxycarbonyl-1,2-cyclohexane, 3,4-epoxycyclohexylmethyl (meth)acrylate or its polymers; and α -olefine epoxides such as epoxyhexadecane.

When the epoxy compound is a resin, its molecular weight ranges between 400 and 100,000, preferably between 400 and 10,000, and the epoxy equivalent weight ranges between 200 and 100,000, preferably between 200 and 5,000.

The epoxy compound may be attached to component (a) or (b) through a covalent bond. This may conveniently be performed by reacting acrylic or methacrylic acid in an amount less than a stoichiometric ratio with an epoxy resin or an epoxy group-containing acrylic resin to produce component (a) having remaining epoxy groups. Component (b) having epoxy groups covalently attached thereto may conveniently be produced by copolymerizing an epoxy group-containing monomer such as glycidyl (meth)acrylate, β -methylglycidyl (meth)acrylate or 3,4-epoxycyclohexylmethyl (meth)acrylate.

Curable resin compositions

The proportions of component (a) and component (b) in the curable resin composition of the present invention generally lie between 2:1 and 1:2, preferably between 1.5:1 and 1:1.5 relative to the double bond and the activated methylene to be added thereto. The proportion of component (c) may range generally between 0.1 and 10.0 equivalent %, preferably between 0.2 and 5.0 equivalent % based on the sum of component (a) and component (b).

All of components (a), (b) and (c) are dissolved or dispersed in an organic solvent conventionally used in the coating industry. Examples thereof include aliphatic hydrocarbons such as hexane or heptane; aromatic hydrocarbons such as toluene or xylene; petroleum fractions having a suitable boiling point range; esters such as butyl acetate, ethylene glycol diacetate or 2-ethylhexyl acetate; ketones such as acetone, methyl ethyl ketone (MEK), methyl isobutyl ketone (MIBK) or cyclohexanone; ethers such as tetrahydrofuran, ethylene glycol dimethyl ether, ethylene glycol diethyl ether or ethylene glycol dibutyl ether; alcohols such as butanol or methoxypropanol; and mixtures of these solvents.

When used as a coating composition, it may contain a variety of conventional additives such as extender pigments, coloring pigments, sag-preventing agents, UV absorbers and the like. After applying onto a substrate as a film, the composition may be cured, depending upon the type of catalyst, by leaving at ambient temperature or baking the film at a temperature between 100 °C and 200°C, preferably between 120°C and 150°C.

The following examples are intended to illustrate the present invention without limiting thereto. All parts and % therein are by weight unless otherwise specified.

0040

~0042

Production Example 1

To a 4-necked flask charged with 360 parts of xylene was added dropwise a monomer mixture containing a polymerization initiator shown in Table 1 at 110°C over 3 hours. After the addition, the mixture was allowed to react at 110°C for 30 minutes. Then 2 parts of KAYAESTER O (t-butylperoxyoctate sold by Nippon Kayaku Co., Ltd.) in 40 parts of xylene were added dropwise at 110 °C over 30 minutes and the mixture allowed to react at the same temperature for additional 90 minutes. An acrylic resin solution having 49.4 % nonvolatiles, a number average molecular weight Mn measured by the GPC method of 5,400, and an activated methylene equivalent weight of 200 was obtained.

Production Examples 2-10

Production Example 1 was followed using a mixture shown in Table 1 and Table 2, respectively. Nonvolatile con-

EP 0 808 860 A2

tents, Mn's and other data are also shown in Table 1 and Table 2.

Table 1

Material, Parts	Production Example				
	1	2	3	4	5
2-(ethoxymalonyloxy)ethyl methacrylate	244	244	244	244	244
n-butyl acrylate	48	129	117	125	129
styrene	80	—	—	—	28
glycidyl methacrylate	28	28	—	—	—
M-100 ¹⁾	—	—	39	—	—
M-GMA ²⁾	—	—	—	31	—
KAYAESTER O	18	18	18	18	18
Nonvolatiles, %	49.4	49.4	50.0	48.7	50.5
Mn	5400	5300	4300	4600	5000
Activated methyle E.W. ³⁾	200	200	200	200	200
Epoxy E.W. ³⁾	2004	2004	2004	2004	—

1) 3,4-epoxycyclohexylmethyl methacrylate, Daicel Chemical Industries, Ltd.

2) β -methylglycidyl methacrylate, Daicel Chemical Industries, Ltd.

3) Resin solids basis.

Table 2

Material, Parts	Production Example				
	6	7	8	9	10
2-(ethoxymalonyloxy)ethyl methacrylate	244	244	244	209	244
n-butyl methacrylate	61	58	61	111	70
styrene	80	80	80	80	80
glycidyl methacrylate	—	—	28	—	—
2-(methacryloyloxy)ethyl trimethylammonium chloride	15	—	—	—	—
2-(methacryloyloxy)ethyl trimethylammonium bromide	—	18	—	—	—
methacrylic acid	—	—	—	—	6
KAYAESTER O	18	18	48	18	18
Nonvolatiles, %	49.9	49.9	49.9	49.9	49.9
Mn	4300	4300	2500	4200	4300
Activated methyle E.W.	200	200	200	468	200
Epoxy E.W.	—	—	2004	—	—

Production Examples 11-15 (for comparison)

Production Example 1 was followed using a mixture shown in Table 3, respectively. Nonvolatile contents, Mn's and other data are shown also in Table 3.

Table 3

Material, parts	Production Example				
	11	12	13	14	15
2-Acetoacetoxyethyl methacrylate	214	214	214	214	214
n-Butyl methacrylate	78	129	129	129	129
Styrene	80	30	30	30	30
Glycidyl methacrylate	28	28	—	—	—
M-100 ¹⁾	—	—	39	—	—
M-GMA ²⁾	—	—	—	31	—
KAYAESTER O	18	18	18	18	18
Nonvolatiles, %	49.4	50.0	51.2	49.6	50.1
Mn	4,900	4,900	4,400	4,600	4,900
Activated methylene E.W. ³⁾	200	200	200	200	200
Epoxy E.W. ³⁾	2,004	2,004	2,004	2,004	—

1) 3,4-epoxycyclohexylmethyl methacrylate, Daicel Chemical Industries, Ltd.

2) β -methylglycidyl methacrylate, Daicel Chemical Industries, Ltd.

3) Resin solids basis.

Example 1

405 parts of the resin solution of Production Example 1 (200 parts as solids), 50 parts of pentaerythritol triacrylate (PETA), 2 parts of tetrabutylammonium bromide (TBABr), 10 parts of UV absorber (TINUBIN 900 sold by Ciba-Geigy AG) and 5 parts of photostabilizer (SANOL LS-400 sold by Ciba-Geigy AG) were thoroughly mixed to make a solution. The solution was applied on a tinplate using #40 bar coater and heated in an oven at 140°C for 20 minutes. The resulting cured film was tested for gel fraction and weatherability. The gel fraction was determined based on the difference of weight before and after soaking the film in a 1:1 mixture of acetone and methanol at room temperature for 48 hours. The weatherability of the cured film was evaluated in terms of degree of yellowing when subjecting to five cycles of standard weathering test in a sunshine weatherometer.

The evaluation schedule is as follows.

Very good (GG): No yellowing.

Good (G): Slightly appreciable yellowing.

Fair (F): Appreciable yellowing.

Bad (B): Remarkable yellowing.

The test results are shown in Table 4.

Examples 2-30

Example 1 was followed, respectively, except that the formulation was changed as shown in Table 4 through Table 9. The results are shown in Table 4 through Table 9.

Table 4

	Example				
	1	2	3	4	5
Component (a), parts	PETA 50	PETA 50	PETA 50	PETA 50	PETA 50
Component (b), parts	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405
Component (c), parts	TBABr 2	TBANO 2	TBAC1 2	TBAPA 2	TBASA 2
Curing condition	140°C x 20min.				
Gel fraction, %	99.8	98.5	97.6	98.1	98.1
Weatherability	GG	GG	GG	GG	GG

PETA: pentaerythritol triacrylate
 TBABr: tetrabutylammonium bromide
 TBANO: tetrabutylammonium nitrate
 TBAC1: tetrabutylammonium chloride
 TBAPA: tetrabutylammonium phthalate
 TBASA: tetrabutylammonium salicylate

Table 5

	Example				
	6	7	8	9	10
Component (a), parts	PETA 50	PETA 50	PETA 50	PETA 50	PETA 50
Component (b), parts	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405
Component (c), parts	TBAB 2	TBAP 2	TBAI 2	TBAAc 2	TMBAC1 2
Curing condition	140°C x 20min.	160°C x 20min.	160°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	98.4	95.2	93.5	98.8	98.1
Weatherability	GG	GG	GG	GG	GG

TBABA: tetrabutylammonium benzoate
 TBAP: tetrabutylammonium di-t-butylphosphate
 TBAI: tetrabutylammonium iodide
 TBAAC: tetrabutylammonium acetate
 TMBAC1: trimethylbenzylammonium chloride

Table 6

	Example				
	11	12	13	14	15
Component (a), parts	TMPTA 50	PETA 50	PETA 50	DPCA-120 150	UA306H 50
Component (b), parts	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.2 405	Pro.Ex.5 396
Component (c), parts	TBPBr 2	TMSI 4	CPBr 4	TBAC1 2	TBABr 2
Epoxy compound parts	—	—	—	—	EPIKOTE 825 15
Curing condition	140°C x 20min.	160°C x 20min.	160°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	98.4	90.2	96.5	92.8	93.1
Weatherability	GG	G	G	GG	GG

TMPTA: trimethylopropane triacrylate
 DPCA-120: KAYARAD DPCA-120, dipentaerythritol/caprolactone (1:12)adduct hexacrylate sold by Nippon Kayaku Co., Ltd.
 UA 306H: HMDI/pentaerythritol (1:2) adduct sold by Kyoeisha Yushi Kagaku Kogyo K.K.
 TBPBr: tetrabutylphosphonium bromide;
 TMSI: trimethylsulfonium iodide;
 CPBr: cetylpyridiniumbromide;
 EPIKOTE 825: bisphenol A diglycidyl ether

Table 7

	Example				
	16	17	18	19	20
Component (a), parts	A-400 200	3002A 200	PETA 50	PETA 50	TMPTA 50
Component (b), parts	Pro.Ex.3 400	Pro.Ex.4 411	Pro.Ex.5 396	Pro.Ex.5 396	Pro.Ex.5 396
Component (c), parts	TBAC1 2	TBAC1 2	TBAOH 2	TBAF 2	NaOMe 2
Curing condition	140°C x 20min.				
Gel fraction, %	93.7	94.8	96.0	94.0	90.4
Weatherability	G	G	GG	GG	G

A-400: NK ESTER A-400, polyethylene glycol diacrylate sold by Shin-Nakamura Kagaku Kogyo K.K.
 3002A: EPOXY ESTER 3002A, reaction product of bisphenol A/propylene oxide (1:2) adduct diglycidyl ether and acrylic acid sold by Kyoeisha Yushi Kagaku Kogyo K.K.
 TBAOH: tetrabutylammonium hydroxide;
 TBAF: tetrabutylammonium fluoride;
 NaOMe: sodium methoxide

Table 8

	Example				
	21	22	23	24	25
Component (a), parts	PETA 50	PETA 50	PETA 50	PETA 50	PETA 50
Component (b), parts	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405	Pro.Ex.1 405
Component (c), parts	Pro.Ex.6 50	TBABH4 2	QDMC 2	TMOBr 2	GTAC 2
Curing condition	140°C x 20min.				
Gel fraction, %	98.9	99.2	97.5	98	97.1
Wetherability	GG	GG	GG	GG	GG

TBABH4: tetrabutylammonium tetrahydroborate
 QDMC: 2-(methacryloyloxy)ethyltrimethylammonium chloride
 TMOBr: trimethyloctylammonium bromide
 GTAC: glycidyltrimethylammonium chloride

Table 9

	Example				
	26	27	28	29	30
Component (a), parts	PETA 50	PETA 50	PETA 50	PETA 50	PETA 50
Component (b), parts	Pro.Ex.1 405	Pro.Ex.8 405	Pro.Ex.9 405	Pro.Ex.10 405	Pro.Ex.10 405
Component (c), parts	Pro.Ex.7 50	TBABr 2	TBABr 2	TBAOH 2	TMBOH 2
Curing condition	140°C x 20min.	140°C x 20min.	140°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	98.2	99.2	97.2	98	97.1
Wetherability	GG	GG	GG	GG	GG

TMBOH: trimethylbenzylammonium hydroxide

Comparative Examples 1-20

Example 1 was followed, respectively, except that the formulation was changed as shown in Table 10 through Table 13. These Comparative Examples correspond to Examples 1-20. The results are shown in Table 10 through Table 13.

Table 10

	Comparative Example				
	1	2	3	4	5
Component (a), parts	PETA 50	PETA 50	PETA 50	PETA 50	PETA 50
Component (b), parts	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405
Component (c), parts	TBABr 2	TBANO 2	TBAC1 2	TBAPA 2	TBASA 2
Curing condition	140°C x 20min.	140°C x 20min.	140°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	98.8	97.8	98.6	98.0	96.9
Weatherability	B	B	B	B	B

Table 11

	Comparative Example				
	6	7	8	9	10
Component (a), parts	PETA 50	PETA 50	PETA 50	PETA 50	PETA 50
Component (b), parts	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405
Component (c), parts	TBABA 2	TBAP 2	TBAI 2	TBAAc 2	TMBAC1 2
Curing condition	140°C x 20min.	160°C x 20min.	160°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	98.8	97.8	98.6	98.0	96.9
Weatherability	B	B	B	B	B

Table 12

	Comparative Example				
	11	12	13	14	15
Component (a), parts	TMPTA 50	PETA 50	PETA 50	DPCA-120 150	UA306H 50
Component (b), parts	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.11 405	Pro.Ex.12 400	Pro.Ex.15 399
Component (c), parts	TBPBr 2	TMSI 4	CPBr 4	TBAC1 2	TBABr 2
Epoxy compound parts	—	—	—	—	EPIKOTE 825 15
Curing condition	140°C x 20min.	160°C x 20min.	160°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	98.7	88.8	86.9	93.8	95.4
Weatherability	B	B	B	B	B

Table 13

	Comparative Example				
	16	17	18	19	20
Component (a), parts	A-400 200	3002A 200	PETA 50	PETA 50	TMPTA 50
Component (b), parts	Pro.Ex.13 391	Pro.Ex.14 403	Pro.Ex.15 399	Pro.Ex.15 399	Pro.Ex.15 399
Component (c), parts	TBAC1 2	TBAC1 2	TBAOH 2	TBAF 2	NaOMe 2
Curing condition	140°C x 20min.	140°C x 20min.	140°C x 20min.	140°C x 20min.	140°C x 20min.
Gel fraction, %	94.5	93.3	96.6	95.8	92.5
Weatherability	B	B	B	B	B

Example 31

55 A zinc phosphate-treated steel plate of 0.8 mm thickness was coated with a cathodic electrodeposition paint (POWER TOP PU-50, Nippon Paint Co., Ltd.) electrically to a dry film thickness of about 25 µm, then with a midlayer paint (ORGANIC P-2 sealer, Nippon Paint Co., Ltd.) by air spraying to a dry film thickness of about 40 µm. Then, both paint films were baked at 140 °C for 30 minutes.

50 In the next step, a base coat composition having the following formulation was air sprayed to a dry film thickness of about 16 µm and allowed to set for about 7 minutes.

Base coat composition

Material	Parts
ALUPASTE 7160N (aluminum flake paste sold by Toyo Aluminum Co., Ltd., flake content 65 %)	10
Acrylic varnish (80 % nonvolatiles, OH #100, acid #80, Mn 1,800)	50
CYMEL 202 (melamine resin sold by Mitsui Cytec Co., Ltd., 80 % nonvolatiles)	10
CYMEL 327 (ditto, 90 % nonvolatiles)	10
Isopropyl alcohol	7

Then, the solution of Example 1 adjusted to Ford cup #4 viscosity of 30 seconds was applied electrostatically onto the base coat film wet-on-wet, and baked both films simultaneously at 140 °C for 25 minutes.

Comparative Example 21

Example 31 was followed except that the solution of Example 1 was replaced with the solution of Comparative Example 1.

Wetherability Evaluation

Cured multilayer coats of Example 31 and Comparative Example 21 were subjected to the wetherability test as in Example 1. The degree of yellowing was not observed with the multilayer Coat of Example 31 while the multilayer coat of Comparative Example 21 was remarkably yellowed.

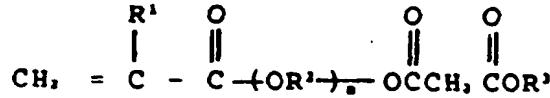
Claims

1. A curable resin composition comprising:

(a) a component containing a plurality of α , β - ethylenically unsaturated carbonyl groups in the molecule;
 (b) an acrylate polymer containing a plurality of malonate-terminated pendant groups in the molecule; and
 (c) a catalyst capable promoting the Michael reaction between said components (a) and (b).

2. The composition of claim 1, wherein said component (a) is a polyol poly(meth)acrylate, an unsaturated polyester resin, a polyester acrylate resin, an epoxy acrylate resin, a urethane acrylate resin, an α , β - unsaturated carbonyl group-containing acrylic resin, a polyether acrylate resin, a silicone oligomer having (meth) acryloyl groups, or an α , β -unsaturated carbonyl group-containing fluororesin.

3. The Composition of claim 1, wherein said acrylate polymer comprises a malonate-terminated acrylate monomer of the formula:



wherein R¹ is hydrogen or methyl, R² is a C₂-C₆ alkylene, R³ is a C₁-C₆ alkyl and n is an integer from 1 to 10 and an ethylenically unsaturated monomer copolymerizable therewith.

4. The composition of claim 3, wherein said malonate-terminated acrylate monomer is an asymmetric malonate ester in which one of carboxyl groups is esterified with an alkanol while the other carboxyl group is esterified with a hydroxyalkyl (meth)acrylate.

5. The composition of claim 3, wherein said malonate-terminated acrylate monomer is an asymmetric malonate ester in which one of carboxyl groups is esterified with an alkanol while the other carboxyl group is esterified with a poly-oxoalkylene glycol mono(meth)acrylate.

6. The Composition of claim 1, wherein said catalyst (c) is a strong base or a quaternary ammonium fluoride or tetrahydroborate.

cl. 7
5 7. The composition of claim 6, wherein said strong base is an alkali metal hydroxide, an alkali metal alkoxide, a quaternary ammonium hydroxide, a tertiary amine, a guanidine, an amidine, a tertiary phosphine, or a mixture thereof.

10 8. The composition of claim 1, wherein said catalyst (c) is a combination of an onium salt with an epoxy compound independently added to the composition or covalently attached to said component (a) or said component (b), said onium salt itself having no catalytic activity on the Michael reaction.

15 9. The composition of claim 1, wherein the onium cation of said onium salt is a tetraalkylammonium, trialkylaralkylammonium, alkylpyridinium, tetraalkylphosphonium or trialkylsulfonium and the counter anion is chloride, bromide, iodide, salicylate, monobasic carboxylate, polybasic carboxylate, nitrate, sulfonate, sulfate, acid sulfate ester, sulfite, phosphate or acid phosphate ester anion.

20 10. The composition of claim 8, wherein said epoxy compound is a glycidyl ether, a glycidyl ester, an alicyclic epoxide or an α -olefin epoxide added to the composition as a discrete component.

11. The composition of claim 8, wherein said epoxy compound is an epoxy function remaining in said component (a), or glycidyl (meth)acrylate, β -methylglycidyl (meth) acrylate or 3,4-epoxycyclohexylmethyl (meth)acrylate copolymerized with the monomers of said acrylate polymer (a).

25 12. The composition of claim 8, wherein said onium salt is a reactive quaternary ammonium salt having a functional group capable of covalently linking to said component (a) or said component (b).

13. The composition of claim 12, wherein said reactive quaternary ammonium salt is a glycidyl ammonium salt independently added to the composition or linked to said component (a) or said component (b) through the reaction with a free carboxyl group thereof.

30 14. The composition of claim 12, wherein said reactive quaternary ammonium salt is an ammonium salt having (meth) acryloyl group independently added to the composition or copolymerized in said component (a) or said component (b).

15. A coating composition comprising the curable resin composition of claim 1 as a vehicle component thereof.

35 16. A coating composition comprising the curable resin composition of claim 12 as a vehicle component thereof.

40

45

50

55



(19) Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) EP 0 808 860 A3

(12) EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
10.02.1999 Bulletin 1999/06

(51) Int. Cl.⁶: C08G 61/12

(43) Date of publication A2:
26.11.1997 Bulletin 1997/48

(21) Application number: 97108237.5

(22) Date of filing: 21.05.1997

(84) Designated Contracting States:
DE FR GB

• Aoki, Kei
Ikoma-shi, Nara-ken (JP)

(30) Priority: 22.05.1996 JP 151546/96

(74) Representative:
Kinzebach, Werner, Dr. et al
Patentanwälte
Reitstötter, Kinzebach und Partner
Postfach 86 06 49
81633 München (DE)

(71) Applicant:
NIPPON PAINT CO., LTD.
Kita-ku, Osaka (JP)

(72) Inventors:
• Irie, Takashi
Suita-shi, Osaka-fu (JP)

(54) Curable resin composition for coating use

(57) A resin composition curable through a Michael reaction between (a) a component having a plurality of α , β -unsaturated carbonyl groups and (b) a component having a plurality of activated methylene group in the presence of a catalyst is disclosed. The activated methylene component is a polymer of an asymmetric malonate ester in which one of carboxyl groups is esterified with an alkanol while the other carboxyl group is esterified with hydroxyalkyl (meth) acrylate or polyoxyalkylene glycol mono(meth)acrylate.



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 97 10 8237

DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
X	US 4 602 061 A (AKKERMANN JAAP M) 22 July 1986 * column 2, line 10 - line 14; claims * * column 3, line 27 - line 65 * ---	1-16	C08F8/46 C08G81/02 C09D133/06 C08G61/12
X	EP 0 596 461 A (HERBERTS & CO GMBH) 11 May 1994 * page 4, column 20-25; claims * * page 9, line 43 - line 54 * ---	1-16	
A	EP 0 227 454 A (COOK PAINT & VARNISH INC) 1 July 1987 * claims; examples * ---	1-16	
TECHNICAL FIELDS SEARCHED (Int.Cl.6)			
C08F C08G C09D C08L			
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	17 December 1998	Friederich, P	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 97 10 8237

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

17-12-1998

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
US 4602061	A	22-07-1986	AU	576837 B	08-09-1988
			AU	4045885 A	03-10-1985
			BR	8501408 A	26-11-1985
			CA	1252240 A	04-04-1989
			DE	3561824 A	14-04-1988
			DK	139485 A , B ,	30-09-1985
			EP	0161697 A	21-11-1985
			FI	851258 A , B	30-09-1985
			GR	850785 A	25-11-1985
			JP	1932175 C	26-05-1995
			JP	6062743 B	17-08-1994
			JP	60223816 A	08-11-1985
EP 0596461	A	11-05-1994	DE	4237492 A	11-05-1994
			AT	164872 T	15-04-1998
			CA	2102170 A	07-05-1994
			DE	59308380 D	14-05-1998
			ES	2115706 T	01-07-1998
			JP	6234954 A	23-08-1994
EP 0227454	A	01-07-1987	BR	8606287 A	06-10-1987
			JP	62223204 A	01-10-1987
			US	4804715 A	14-02-1989

THIS PAGE BLANK (USPTO)